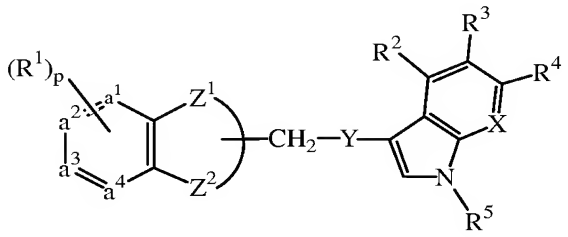
This listing of claims will replace all prior versions, and listings, of claims in the application.

## Listing of Claims:

1. (Currently Amended) A compound according to Formula (I)



(I)

a pharmaceutically acceptable acid or base addition salt thereof, a stereochemically isomeric form thereof, an N-oxide form thereof or a quaternary ammonium salt thereof, wherein

$-a^1=a^2-a^3=a^4-$ is a bivalent radical of formula

~~-N=CH-CH=CH- (a-1),~~

~~-CH=N-CH=CH- (a-2),~~

-CH=CH-N=CH- (a-3) or

-CH=CH-CH=N- (a-4) ;

$-Z^1-Z^2-$ is a bivalent radical of formula

$-O-CH_2-O-$ (b-1),

$-O-CH_2-CH_2-O-$ (b-2),

$-NR^7-CH_2-CH_2-O-$ (b-3),

~~$-O-CH_2-CH_2-NR^7-$ (b-4),~~

~~$-NR^7-CH_2-CH_2-NR^7-$ (b-5) or~~

~~-S-CH2-CH2-O- (b-6)~~ ;

wherein $R^7$ is hydrogen, hydroxy, alkyl, alkyloxyalkyl or alkylcarbonyl ;

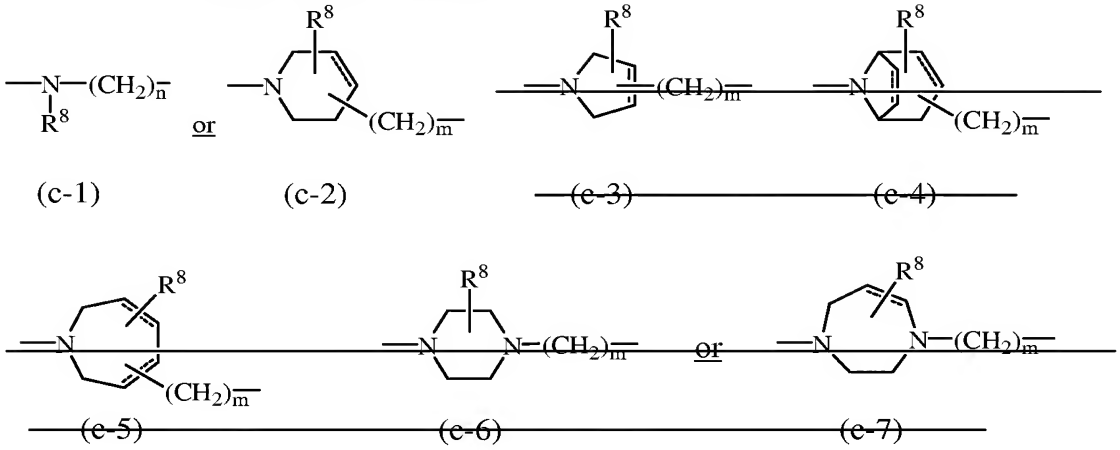
X is $CR^6$ ~~or N~~;

each $R^1$, $R^2$, $R^3$, $R^4$ and $R^6$ is independently hydrogen, halo, cyano, nitro, <u>or</u> ~~alkyl, alkenyl, mono- or dialkylaminoalkyl,~~ hydroxy~~, alkyloxy, alkylcarbonyloxy, amino, mono- or dialkylamino, formylamino, alkylcarbonylamino, alkylsulfonylamino, hydroxycarbonyl, alkyloxycarbonyl, aminocarbonyl, mono- or dialkylaminocarbonyl, alkylcarbonyloxy-alkyloxycarbonyloxy, alkylthio, aryl or heteroaryl~~;

p is an integer equal to 0, 1, 2 or 3 ;

$R^5$ is hydrogen or alkyl ;

Y is a bivalent radical of formula



(c-1) (c-2) ~~(c-3)~~ ~~(c-4)~~

~~(c-5)~~ ~~(c-6)~~ ~~(c-7)~~

wherein

m is an integer equal to 0 or 1 ;

n is an integer equal to 0, 1, 2, 3, 4, 5 or 6 ;

the dotted line represents an optional double bond ;

$R^8$ is hydrogen~~, halo, alkyl, hydroxy, alkyloxy, alkylcarbonyloxy, alkyloxycarbonyloxy, hydroxycarbonyl, aminocarbonyl, mono- or dialkylaminocarbonyl, alkyloxycarbonyl or amino~~;

alkyl represents a straight or branched saturated hydrocarbon radical having from 1 to 6 carbon atoms or a cyclic saturated hydrocarbon radical having from 3 to 6 carbon atoms; said radical being optionally substituted with at least one phenyl, halo, cyano, oxo, hydroxy, formyl or amino radical;

alkenyl represents a straight or branched unsaturated hydrocarbon radical having from 1 to 6 carbon atoms or a cyclic unsaturated hydrocarbon radical having from 3 to 6 carbon atoms ; said radical having at least one-double bond and said radical being optionally substituted with at least one-phenyl, halo, cyano, oxo, hydroxy, formyl or amino radical;

aryl represents phenyl or naphthyl, optionally substituted with at least one-radical that is alkyl, halo, cyano, oxo, hydroxy, alkyloxy or amino ; and

heteroaryl is a monocyclic heterocyclic radical that is azetidinyl, pyrrolidinyl, dioxolyl, imidazolidinyl, pyrrazolidinyl, piperidinyl, homopiperidinyl, dioxyl, morpholinyl, dithianyl, thiomorpholinyl, piperazinyl, imidazolidinyl, tetrahydrofuranyl, 2H-pyrrolyl, pyrrolinyl, imidazolinyl, pyrrazolinyl, pyrrolyl, imidazolyl, pyrazolyl, triazolyl, furanyl, thienyl,

oxazolyl, isoxazolyl, thiazolyl, thiadiazolyl, isothiazolyl, pyridinyl, pyrimidinyl, pyrazinyl, pyridazinyl or triazinyl ; each radical optionally substituted with at least one-radical that is alkyl, aryl, arylalkyl, halo, cyano, oxo, hydroxy, alkyloxy or amino;

with the proviso that compounds wherein simultaneously -$a^1$=$a^2$-$a^3$=$a^4$- is (a-4), -$Z^1$——$Z^2$- is (b-2) and Y is (c-2) are excluded.

2. (Canceled)

3. (Currently Amended) The compound according to claim 1, ~~wherein $Z^1$-$Z^2$- is a bivalent radical of formula (b-1), (b-2) or (b-3)~~ wherein $R^7$ is hydrogen or methyl.

4. (Previously Presented) The compound according to claim 1, wherein Y is a bivalent radical of formula (c-1) wherein n = 3 and $R^8$ is hydrogen or of formula (c-2) wherein m = 0 or 1 and $R^8$ is hydrogen.

5. (Previously Presented) The compound according to claim 1, wherein X is $CR^6$; $R^2$, $R^3$, $R^4$ and $R^6$ are each independently hydrogen, halo, cyano, nitro or hydroxy; and $R^5$ is hydrogen.

6. (Currently Amended) The compound according to claim 1, ~~wherein $a^1$=$a^2$-$a^3$=$a^4$- is a bivalent radical of formula (a-3) or (a-4) ; $Z^1$-$Z^2$- is a bivalent radical of formula (b-1), (b-2) or (b-3)~~ wherein $R^7$ is hydrogen or methyl; Y is a bivalent radical of formula (c-1) wherein n = 3 and $R^8$ is hydrogen or (c-2) wherein ~~m = 0 or 1 and~~ $R^8$ is hydrogen; ~~X is $CR^6$; $R^2$, $R^3$, $R^4$ and $R^6$ are each independently hydrogen, halo, cyano, nitro or hydroxy~~ and $R^5$ is hydrogen.

7. (Canceled)

8. (Previously Presented) A pharmaceutical composition comprising a pharmaceutically acceptable carrier or diluent and, as active ingredient, a therapeutically effective amount of a compound according to claim 1.

9. (Withdrawn) A method for the prevention and/or treatment in a mammal of a disorder or disease responsive to the inhibition of dopamine $D_2$, $D_3$ and/or $D_4$-receptors,

comprising administering to the mammal a therapeutically effective amount of a compound according to claim 1.

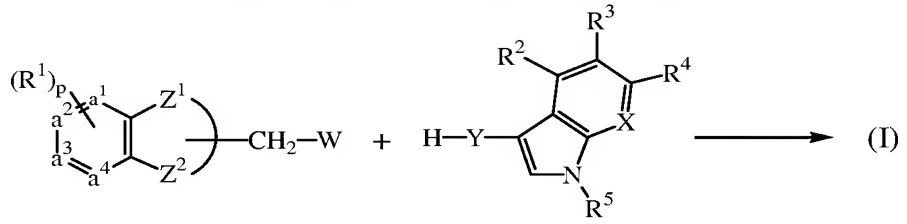
10. (Withdrawn) A method for the prevention and/or treatment in a mammal of a disorder or disease responsive to the inhibition of serotonin reuptake and antagonism of 5-$HT_{1A}$ receptors, comprising administering to the mammal a therapeutically effective amount of a compound according to claim 1.

11. (Withdrawn) A method for the prevention and/or treatment in a mammal of a disorder or disease responsive to the combined effect of a dopamine $D_2$, $D_3$ and/or $D_4$ antagonist, a selective serotonin reuptake inhibitor (SSRI) and a 5-$HT_{1A}$-agonist, partial agonist or antagonist, comprising administering to the mammal a therapeutically effective amount of a compound according to claim 1.

12. (Withdrawn) A method for the prevention and/or treatment in a mammal of general anxiety disorder, panic disorder, obsessive compulsive disorder, depression, social phobia, eating disorders, psychosis or neurological disorders, comprising administering to the mammal a therapeutically effective amount of a compound according to claim 1.

13. (Withdrawn) A method for the prevention and/or treatment of schizophrenia in a mammal, comprising administering to the mammal a therapeutically effective amount of a compound according to claim 1.

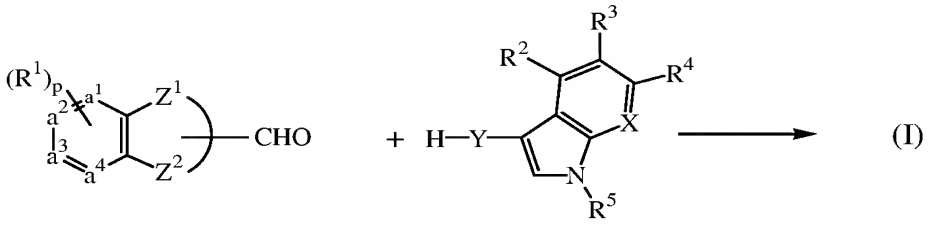
14. (Withdrawn/Currently Amended) A process for the preparation of a compound according to Formula (I) comprising
--alkylating a compound-of Formula (III) with a compound of Formula (II), , in a reaction-inert solvent and optionally in the presence of a base;



(II) (III)

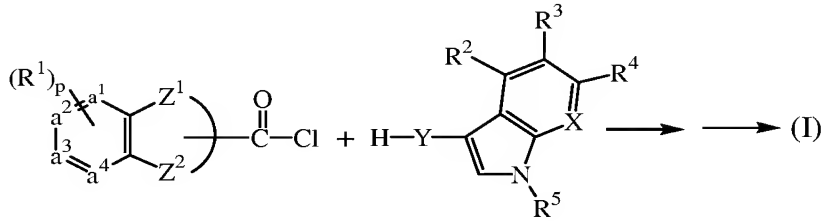
wherein W is a leaving group; or

--reductively aminating a compound-of Formula (IV) ~~is~~ with a compound-of Formula (III) in a reaction-inert solvent and in the presence of a reducing agent; or



(IV) (III)

--reacting an acid chloride of Formula (V) with a compound-of Formula (III) in a reaction-inert solvent and in the presence of a suitable base, and reducing the amide in a reaction-inert solvent in the presence of a reducing agent;



(V) (III)

$-a^1=a^2-a^3=a^4-$ is a bivalent radical of formula

~~-N=CH-CH=CH- (a-1),~~

~~-CH=N-CH=CH- (a-2),~~

-CH=CH-N=CH- (a-3) or

-CH=CH-CH=N- (a-4) ;

$-Z^1$——$Z^2-$ is a bivalent radical of formula

-O-$CH_2$-O- (b-1),

-O-$CH_2$-$CH_2$-O- (b-2),

-$NR^7$-$CH_2$-$CH_2$-O- (b-3),

~~-O-$CH_2$-$CH_2$-$NR^7$- (b-4),~~

~~-$NR^7$-$CH_2$-$CH_2$-$NR^7$- (b-5) or~~

~~-S-CH2-$CH_2$-O- (b-6)~~ ;

wherein $R^7$ is hydrogen, hydroxy, alkyl, alkyloxyalkyl or alkylcarbonyl;

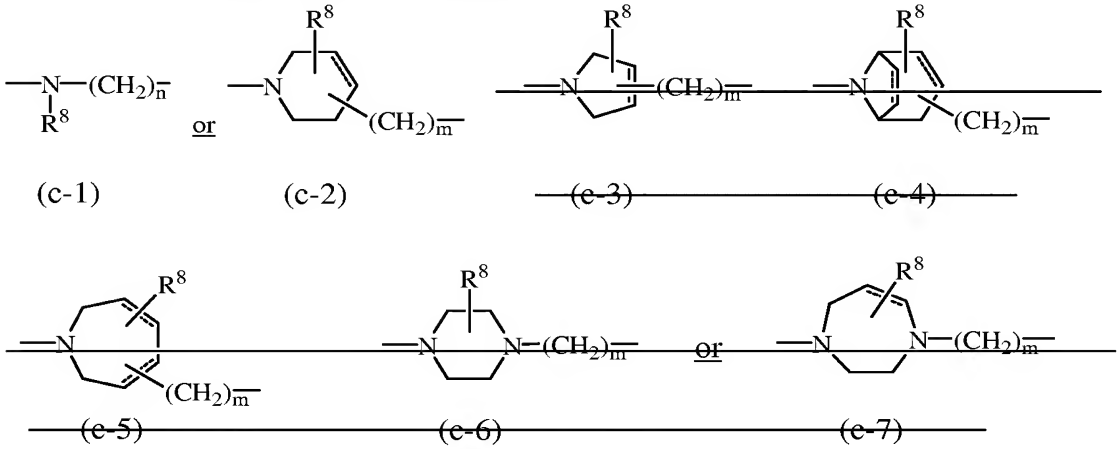
X is $CR^6$ ~~or N~~;

each $R^1$, $R^2$, $R^3$, $R^4$ and $R^6$ is independently hydrogen, halo, cyano, nitro, or ~~alkyl, alkenyl, mono- or dialkylaminoalkyl,~~ hydroxy~~, alkyloxy, alkylcarbonyloxy, amino, mono- or dialkylamino, formylamino, alkylcarbonylamino, alkylsulfonylamino, hydroxycarbonyl, alkyloxycarbonyl, aminocarbonyl, mono- or dialkylaminocarbonyl, alkylcarbonyloxy alkyloxycarbonyloxy, alkylthio, aryl or heteroaryl~~;

p is an integer equal to 0, 1, 2 or 3 ;

$R^5$ is hydrogen or alkyl ;

Y is a bivalent radical of formula



(c-1) or (c-2) ~~(c-3) (c-4)~~

~~(c-5) (c-6)~~ or (c-7)

wherein

m is an integer equal to 0 or 1 ;

n is an integer equal to 0, 1, 2, 3, 4, 5 or 6 ;

the dotted line represents an optional double bond ;

$R^8$ is hydrogen~~, halo, alkyl, hydroxy, alkyloxy, alkylcarbonyloxy, alkyloxycarbonyloxy, hydroxycarbonyl, aminocarbonyl, mono- or dialkylaminocarbonyl, alkyloxycarbonyl or amino~~;

alkyl represents a straight or branched saturated hydrocarbon radical having from 1 to 6 carbon atoms or a cyclic saturated hydrocarbon radical having from 3 to 6 carbon atoms; said radical being optionally substituted with at least one phenyl, halo, cyano, oxo, hydroxy, formyl or amino radical;

alkenyl represents a straight or branched unsaturated hydrocarbon radical having from 1 to 6 carbon atoms or a cyclic unsaturated hydrocarbon radical having from 3 to 6 carbon atoms ; said radical having at least one-double bond and said radical being optionally substituted with at least one-phenyl, halo, cyano, oxo, hydroxy, formyl or amino radical;

aryl represents phenyl or naphthyl, optionally substituted with at least one-radical that is alkyl, halo, cyano, oxo, hydroxy, alkyloxy or amino ; and

heteroaryl is a monocyclic heterocyclic radical that is azetidinyl, pyrrolidinyl, dioxolyl, imidazolidinyl, pyrrazolidinyl, piperidinyl, homopiperidinyl, dioxyl, morpholinyl, dithianyl, thiomorpholinyl, piperazinyl, imidazolidinyl, tetrahydrofuranyl, 2H-pyrrolyl, pyrrolinyl, imidazolinyl, pyrrazolinyl, pyrrolyl, imidazolyl, pyrazolyl, triazolyl, furanyl, thienyl, oxazolyl, isoxazolyl, thiazolyl, thiadiazolyl, isothiazolyl, pyridinyl, pyrimidinyl, pyrazinyl, pyridazinyl or triazinyl ; each radical optionally substituted with at least one-radical that is alkyl, aryl, arylalkyl, halo, cyano, oxo, hydroxy, alkyloxy or amino.

15. (Withdrawn) The process of claim 14, further comprising converting the compound of Formula (I) into a therapeutically active, non-toxic acid addition salt by treatment with an acid.

16. (Withdrawn) The process of claim 15, further comprising converting the acid addition salt into a free base by treatment with alkali.

17. (Withdrawn) The process of claim 16, further comprising converting the compound of Formula (I) into a stereochemically isomeric form, a N-oxide, or a quaternary ammonium salt.

18. (Withdrawn) The process of claim 14, further comprising converting the compound of Formula (I) into a therapeutically active, non-toxic base addition salt by treatment with a base.

19. (Withdrawn) The process of claim 18, further comprising converting the base addition salt into a free acid by treatment with an acid.